



Improving models of species ecological niches: a remote sensing overview

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Abstract: Effective conservation capable of mitigating global biodiversity declines require thorough knowledge on species distributions and their drivers. A species ecological niche determines its geographic distribution, and species distribution models (SDMs) can be used to predict them. For various reasons, e.g., the lack of spatial data on relevant environmental factors, SDMs fail to characterize important ecological relationships. We argue that SDMs do not yet include relevant environmental information, which can be measured with remote sensing (RS). RS may benefit SDMs because it provides information on e.g., ecosystem function, health and structure, complete spatial assessment, and reasonable temporal repeat for the processes that determine geographical distributions. However, RS data is still seldom included in such studies with the exception of climate data. Here we provide a guide for researchers aiming to improve their SDM studies, describing how they might include RS data in their specific study. We propose how to improve models of species ecological niches, by including measures of habitat quality (e.g., productivity), nutritional values, and seasonal or life-cycle events. To date, several studies have shown that using ecologically-relevant environmental predictors derived from RS improve model performance and transferability, and better approximate a species ecological niche. These data, however, are not a panacea for SDMs, as there are cases in which RS predictors are not appropriate, too costly, or exhibit low predictive power. The integration of multiple environmental predictors derived from RS in SDMs can thus improve our knowledge on processes driving biodiversity change and improve our capacity for biodiversity conservation.

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Improving Models of Species Ecological Niches: A Remote Sensing Overview

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Effective conservation capable of mitigating global biodiversity declines require thorough knowledge on species distributions and their drivers. A species ecological niche determines its geographic distribution, and species distribution models (SDMs) can be used to predict them. For various reasons, e.g., the lack of spatial data on relevant environmental factors, SDMs fail to characterize important ecological relationships. We argue that SDMs do not yet include relevant environmental information, which can be measured with remote sensing (RS). RS may benefit SDMs because it provides information on e.g., ecosystem function, health and structure, complete spatial assessment, and reasonable temporal repeat for the processes that determine geographical distributions. However, RS data is still seldom included in such studies with the exception of climate data. Here we provide a guide for researchers aiming to improve their SDM studies, describing how they might include RS data in their specific study. We propose how to improve models of species ecological niches, by including measures of habitat quality (e.g., productivity), nutritional values, and seasonal or life-cycle events. To date, several studies have shown that using ecologically-relevant environmental predictors derived from RS improve model performance and transferability, and better approximate a species ecological niche. These data, however, are not a panacea for SDMs, as there are cases in which RS predictors are not appropriate, too costly, or exhibit low predictive power. The integration of multiple environmental predictors derived from RS in SDMs can thus improve our knowledge on processes driving biodiversity change and improve our capacity for biodiversity conservation.

Keywords: ecological niche, species conservation, remote sensing, species distribution (niche) model, ecological theory

Global environmental change drives biodiversity changes and declines, and the rearrangement of biotic communities (Pereira et al., 2012; Dirzo et al., 2014). The effects of biodiversity declines may lead to losses in the integrity and functioning of ecosystems and the services they provide, thus potentially putting human well-being at risk (Cardinale et al., 2012). International mitigation programs were established to systematically monitor spatial patterns of species distributions (Schmeller et al., 2015). Species distribution models (SDMs) are empirical statistical approaches that use environmental predictors to estimate the species niche, and from there extrapolate the

extent of that niche in space to predict the species distribution (Guisan and Zimmermann, 2000), and have become an important tool in conservation science, planning, and management (Elith and Leathwick, 2009; Pecl et al., 2017).

The choice of environmental predictors is fundamental for SDMs; yet, such selection still remains a main source of debate (Synes and Osborne, 2011). Predictors should measure the processes that link environmental conditions to species occurrence, and match the spatial and temporal scales at which such processes occur (Lechner et al., 2012). Therefore, ecologically relevant predictors are capable of generating robust and models transferable to other regions, time periods or conditions. This is particularly relevant when these models are to be used as a base for conservation planning, considering the effects of global change. Predictors may also differ across taxonomic groups, for example soil type might be good predictors for plants, while forest fragmentation, or temperature might be good predictors for animals (Bradley et al., 2012).

Remote sensing (RS) data allow measuring vegetation condition (Turner et al., 2003), ecosystem productivity (Running et al., 2004), seasonality (Reed et al., 1994), all of which might be used in SDMs (He et al., 2015). These measurements are now available over time series (e.g., Landsat time series; Kennedy et al., 2014), thus expanding the possibilities to model species distributions over time. Upcoming sensors are expected to provide even better and more diverse measurements at finer spatial, temporal and spectral resolutions (e.g., Sentinel satellites; Berger and Aschbacher, 2012). The integration of such remotely sensed information in SDMs can lead to global mapping of biodiversity change (Ferrier, 2011), and thus aim at effective conservation actions (Rose et al., 2015).

Although the use of RS in SDMs is widely advocated and applied (Bradley and Fleishman, 2008; Cord et al., 2013), we argue that it's yet to be explored to its full potential. Since the publication of these studies both RS and SDM science has advanced substantially. On the RS side there have been major developments over the last 10 years, namely the continuation of missions and sensors for multispectral data (Sentinel, Landsat 8), novel sensors that provide additional data on ecosystem functioning which could explain geographic distribution patterns (e.g., Flex; Coppo et al., 2017), and the test studies for other upcoming missions (e.g., Guanter et al., 2015; Lee et al., 2015; Stavros et al., 2017; EnMAP, HypIRI, GEDI). Further data became more easily accessible, in analysis-ready products, and with higher temporal frequency. On the SDM side new algorithms were developed, algorithms were tested for performance and tools were developed to aid researchers on algorithm selection, as well as on getting a better understanding of the ecological meaning of model outputs. Also on the biological data side, many more records were digitized, there was an emergence of citizen science platforms to collect data, and global analyses for multiple taxonomic groups became possible. In this paper we present our perspective on these advances and provide examples on how the last 10 years of RS and SDM have merged and where could they go into the future.

Improving Models of Species Ecological Niches

Soberón (2007) proposes that, to realistically reflect the ecology of a species and the spatial scale at which different processes occur, the model that describes its niche should have abiotic variables depicted at low spatial resolution and biotic variables at high spatial resolution, both interacting dynamically. Thus selecting the best RS predictors to model animal species distribution varies if the goal is to understand how abiotic interactions determine a species niche (Grinnellian niche) or how biotic interactions influence it (Eltonian niche). There is growing evidence that land cover classes generally lead to models unsuitable for prediction (Cord et al., 2014), particularly so when used as proxies for habitat quality or resource availability (Bradley and Fleishman, 2008; Vallecillo et al., 2009), although this also depends on the thematic detail of the land cover maps (Cord et al., 2014). Since Bradley and Fleishman (2008), several studies were conducted that took advantage of the existing RS data. Novel approaches have shown progress toward understanding species responses to abiotic conditions such as climate (Austin and van Niel, 2011), nutritional value (Sheppard et al., 2007) or food resources (Coops et al., 2009), and seasonal variation (Leitão et al., 2010).

Several RS datasets are currently available that could measure several niche axes of species, namely: (i) habitat quality—condition of a habitat type, (ii) nutritional value—food resources available, and (iii) seasonality and life-cycle—temporal variability in habitat due to seasons or individuals, populations and species life-cycles (Gounand et al., 2018). We chose these three niche axes because they capture the most commonly studied aspects of species distribution models and can be more directly measured by RS. In **Table 1** we provide examples that either illustrate instances where RS variables have been used to describe these niche axes in animal SDMs studies, or literature where these variables have been suggested as good proxies to do so. Where we are unaware of a study that uses RS variables, we propose a variable (set of variables) and provide the reference describing the variable itself.

Uncritical use of categorical predictors in SDMs has been discouraged (Elith and Leathwick, 2009). Instead, predictors such as vegetation condition and properties (Zimmermann et al., 2007), phenology (Osborne et al., 2001; Leitão et al., 2010) or structure (Bradbury et al., 2005) could be more informative because they may explain availability of resources, shelter, etc. (Coops et al., 2009; Santos et al., 2016). These, together with databases like e.g., PanTHERIA (Jones et al., 2009), COMADRE (Salguero-Gómez et al., 2016), or COMPADRE (Salguero-Gómez et al., 2015) that include a description of species traits could provide good links to what constitutes a species habitat. Adding information on habitat fragmentation could also be an improvement; and there are many metrics available in the ecological literature (Moilanen and Hanski, 2001; Kindlmann and Burel, 2008). Abiotic climatic data can explain a species distribution because climate parameters like temperature and precipitation directly drive physiological processes, such as thermoregulation, and thus affect a species geographic distribution (Kearney and Porter, 2009). Also, given that

TABLE 1 | Potential RS variables for describing three axes of a species ecological niche: habitat quality, nutritional value, and seasonality and life cycle.

Environmental drivers	RS predictors	Habitat quality	Nutritional value	Seasonality/life cycle	Selected references
SOIL					
Soil type	Spectral features*, such as reflectance in the absorption region of specific constituent minerals, etc.	✓	✓		Guanter et al., 2015
Soil moisture	Spectral indices* (e.g., NDWI) or transformations (e.g., wetness); data from the SMOS Earth Explorer	✓	✓		Papes et al., 2012
CLIMATE					
Temperature	Thermal data* (LST)	✓		✓	Cord and Rödger, 2011
Precipitation	Cloud cover*; precipitation data derived from CHIRPS	✓	✓	✓	Wilson and Jetz, 2016
VEGETATION					
Vegetation structure	Laser scanning metrics* (e.g., tree height, canopy height, canopy vertical structure, etc.); parameters derived from RTM	✓	✓		Bradbury et al., 2005
Vegetation condition	Spectral indices* (e.g., NDVI, EVI) or transformations (greenness and brightness); parameters derived from RTM	✓	✓		Santos et al., 2016
Productivity	Biophysical parameters* (e.g., fPAR, LAI); parameters derived from RTM	✓	✓	✓	Coops et al., 2009
Plant stress	Spectral indices* (e.g., PRI, EWT); fluorescence data	✓	✓	✓	Saatchi et al., 2008
Land surface phenology	Phenological metrics from time series* (e.g., start/length of the growing season, senescence, etc.)	✓	✓	✓	Leitão et al., 2010
Nutrients	Spectral features*, such as reflectance in specific absorption features of nitrogen, etc.	✓	✓		Sheppard et al., 2007
Landscape configuration	Landscape and surface metrics relating to fragmentation, connectivity, heterogeneity, texture*, etc.	✓		✓	Bellis et al., 2008
Habitat information	Habitat type (LCC), fractional cover* of functional types (trees, grass, etc.)	✓	✓	✓	Wessels et al., 2004
DISTURBANCES					
Disturbances	Distance metrics* (e.g., to nearest road or settlement); Change products from LCC or fractional cover; Indices derived from time series (e.g., DI)	✓			Devictor et al., 2008
Human Impact	Stable nighttime lights* derived from the DMSP, land use intensity	✓			Escobar et al., 2015

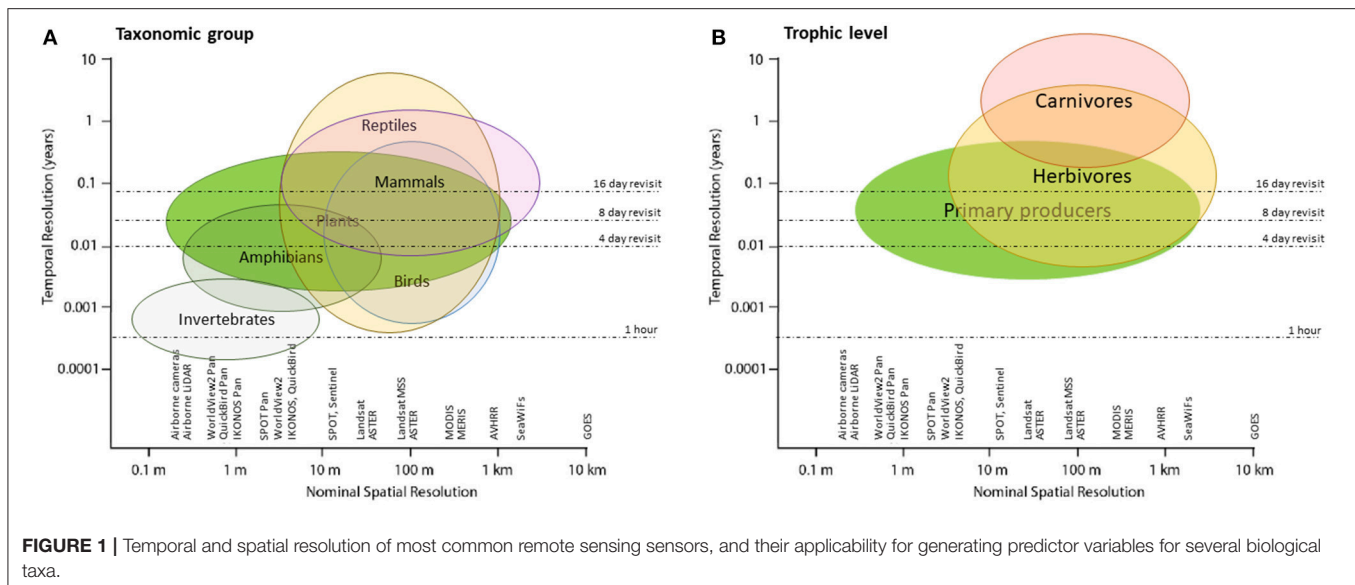
NDWI, Normalized Difference Water Index; SMOS, Soil Moisture Ocean Salinity; LST, Land Surface Temperature; CHIRPS, Climate Hazards group InfraRed Precipitation with Station data; RTM, Radiative Transfer Models; NDVI, Normalized Difference Vegetation Index; EVI, Enhanced Vegetation Index; fPAR, fraction of Photosynthetically Active Radiation; LAI, Leaf Area Index; PRI, Photochemical Reflectance Index; EWT, Equivalent Water Thickness; LCC, Land Cover Classification; DI, Disturbance Index; DMSP, Defense Meteorological Satellite Program. *Denotes which variable is used in the selected reference.

predictors that more directly reflect the species ecology generate transferable models (Austin, 2002), historical reconstructions or extreme climatic events may provide unique real-life experiments and opportunities to test the utility of predictors for model transferability (Randin et al., 2006; Santos et al., 2017). All these approaches can be applicable to RS predictors and integrated in SDMs (Bellis et al., 2008).

In order to characterize *habitat quality* of a species it is relevant to e.g., explore spectral indices that measure functional aspects of ecosystems (Ustin et al., 2004; Pettorelli et al., 2018), such as the depiction of productivity (e.g., through the Normalized Difference Vegetation Index, NDVI; Santos et al.,

2016). Other properties like canopy water condition and soil properties may also be measured by indices (Table 1). If one is interested in adding information on vegetation structure, active RS products like active airborne laser scanners (LiDAR) can be an option (Vierling et al., 2008). LiDAR data can be used to measure the vertical structure of vegetation, while satellite RADAR data (from RADARSAT-1 and Sentinel-1) can retrieve vegetation morphology or elevation. Vegetation structural measures can be used directly used as predictors of suitable habitat for wildlife species (Bradbury et al., 2005).

If the goal is to understand how *nutritional quality* affects animal species distributions, then it is important to consider the



trophic level of the studied species. For example, RS measures of productivity and pigment concentration may be indicators of nutritional quality of plants (Field et al., 1995) or fruiting events for herbivores (Sudbrink et al., 2003). Productivity may also be approximated by fPAR, although forthcoming sensors will be able to measure photosynthetic activity directly. Both of these metrics could be a good proxy of nutritional value for herbivores/granivores (Leitão et al., 2010). For carnivores, productivity might indicate where their prey is and local studies have demonstrated that it improves model performance (Santos et al., 2016).

Finally, if one is interested in how species distributions may vary *seasonally and within a life cycle*, time series data on fPAR, vegetation fractional cover, NDVI and Enhanced Vegetation Index (EVI), etc. are becoming more readily available (Bischof et al., 2012). For example, migratory species respond to the greening of the vegetation which determines their migratory patterns (Post et al., 2003; Bischof et al., 2012). Cord and Rödder (2011) demonstrated that the use of multi-temporal RS predictors (EVI and Land Surface Temperature—LST—from MODIS) improved the modeling of the distributions of eight Mexican amphibian species with differing habitat preferences. This is a very promising avenue as it allows modeling changes in species distributions over time, and assessing how vegetation phenology changes might affect habitat quality and migratory and dispersal processes (Post et al., 2003).

Perspectives for Use of RS in SDMs

RS predictors should not be perceived as a panacea for SDMs, as there are cases where such predictors are not appropriate, too costly, or of low predictive power. In the case of species with small ranges (e.g., amphibians with restricted distribution in mountain tops) satellite data may not provide good coverage, and acquisition and processing of airborne or drone hyperspectral or laser scanning data might be too costly. Models for different

species likely require different predictors, for example, NDVI might be a good predictor for herbivores (Kuemmerle et al., 2014), but not for carnivores (Bradley et al., 2012). However, for plants the choice of NDVI as a predictor is contested, some authors argue that it should not be included as it is correlated with plant productivity (Bradley et al., 2012), while others argue that it is a measure of ecosystem functioning and therefore indicates the relation between productivity and biodiversity. Further a RS image is a snapshot and may not always precisely capture the variability in niche conditions of a species. RS predictors tend to be highly correlated and depending on the modeling algorithm used, it may pick the “wrong” ones that lack ecological significance (Dormann et al., 2013).

There are trade-offs between RS systems, as high resolution in one dimension often comes at the cost of other dimensions; for example, high temporally resolved MODIS data come at a coarse spatial and moderate spectral resolution. It is also important to acknowledge that an index from one sensor may not correspond exactly to that same index from another sensor (see Figure 1). For example, a model calibrated from one sensor’s NDVI is not easily applied to NDVI from another sensor (Roy et al., 2016). Efforts exist within the RS community to calibrate indices across sensors for wider use (Roy et al., 2016), prompted by the recent increase in availability of free and open access satellite imagery (e.g., Sentinel, Landsat, SPOT). The next generation sensors have reheated the effort toward harmonization and intrinsic transferability of multi-sensor data (Wulder et al., 2015). Similarly, consistency in time series within the same sensor needs to be properly calibrated (van Leeuwen et al., 2006). Achieving consistent environmental predictors would potentially allow a generalized and transferable use and development of SDMs.

Novel RS products increase the array of possible predictors to use in SDMs. Novel off-the-shelf RS products which relate to ecosystem processes illustrate well an increasing communication between what ecologists need and what the RS community can provide (Buchanan et al., 2015). Space agencies and

satellite data providers increasingly deliver higher-level products for use in SDMs and therefore minimize the requirement of users to know the details about data processing. Recent and forthcoming satellites, for example hyperspectral sensors such as the Environmental Mapping and Analysis Program (EnMAP) HyperSpectral Imager (HSI; Guanter et al., 2015) or the Hyperspectral InfraRed Imager (HyspIRI; Lee et al., 2015) will allow novel ways to measure physiological properties of vegetation, such as water stress, cellulose or nitrogen contents, or nutrient stress (Leitão et al., 2015). New sensors for chlorophyll fluorescence, such as FLORIS onboard the planned FLEX mission (Coppo et al., 2017), will be able to measure photosynthetic activity directly. The planned BIOMASS radar mission (Le Toan et al., 2011) aims at mapping forest biomass on a global scale.

SDM's themselves include a set of limitations, such as assuming equilibrium between the species and the environment, failing to include the potential for evolutionary adaptation, omitting species interactions and non-climatic constraints, failing to allow for novel climates or conditions, etc. (Schwartz, 2012), but many of such limitations are beyond our proposal to better integrate RS in SDM. In fact, RS can solve some of the issues and others are SDM specific, and both disciplines should converge in some aspects. For example, here we argue that RS could provide a means to include non-climatic distribution constraints, such as habitat or human impact (e.g., land use intensity; Kleijn et al., 2009).

RS data are well suited to integration in SDMs to predict the response of biodiversity to environmental drivers and provide information to intergovernmental forums like the Intergovernmental Panel on Climate Change (IPCC) and the Intergovernmental Science-Policy Platform on Biodiversity and

Ecosystem Services (IPBES). RS for biodiversity has become a priority and has been acknowledged by many international efforts, e.g., Aichi Biodiversity Targets, Sustainable Development Goals, Group on Earth Observation Biodiversity Observation Network, and Essential Biodiversity Variables (Pereira et al., 2013; Skidmore et al., 2015; Pettoirelli et al., 2016). These are all strong arguments for accepting the challenge of finding a set of ecologically relevant RS predictors that may be used to globally understand biodiversity change (Ferrier, 2011; Jetz et al., 2016).

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All authors listed have made a substantial, direct and intellectual contribution to the work, and approved it for publication.

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Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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